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EUV, XUV, and X-RAY Wavelength Sources Created From Laser Plasma Produced From Liquid Metal Solutions

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Richardson

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(45) **Date of Patent:** **Dec. 14, 2004**

(54) **EUV, XUV, AND X-RAY WAVELENGTH
SOURCES CREATED FROM LASER
PLASMA PRODUCED FROM LIQUID
METAL SOLUTIONS**

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patent is extended or adjusted under 35
U.S.C. 154(b) by 0 days.

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2000.

(51) **Int. Cl.**⁷ **H01J 35/08**

(52) **U.S. Cl.** **378/119; 378/143**

(58) **Field of Search** **378/119, 143,**
378/34; 372/5

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Primary Examiner—David V. Bruce

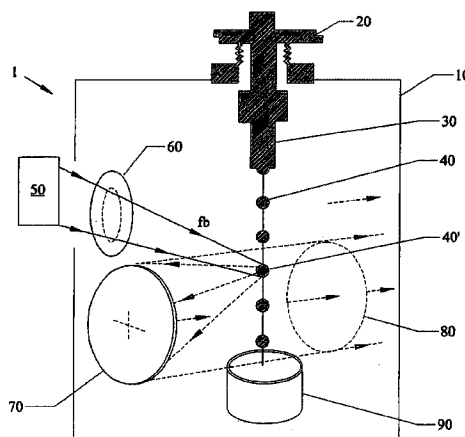
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(57) **ABSTRACT**

Metallic solutions at room temperature used a laser point source target droplets. Using the target metallic solutions results in damage free use to surrounding optical components since no debris are formed. The metallic solutions can produce plasma emissions in the X-rays, XUV, and EUV (extreme ultra violet) spectral ranges of approximately 11.7 nm and 13 nm. The metallic solutions can include molecular liquids or mixtures of elemental and molecular liquids, such as metallic chloride solutions, metallic bromide solutions, metallic sulphate solutions, metallic nitrate solutions, and organo-metallic solutions. The metallic solutions do not need to be heated since they are in a solution form at room temperatures.

74 Claims, 5 Drawing Sheets



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Fig. 1a
(Prior Art)

Spectra of Copper (Cu) target irradiated under similar
Scale: note 10Å = 1 nm

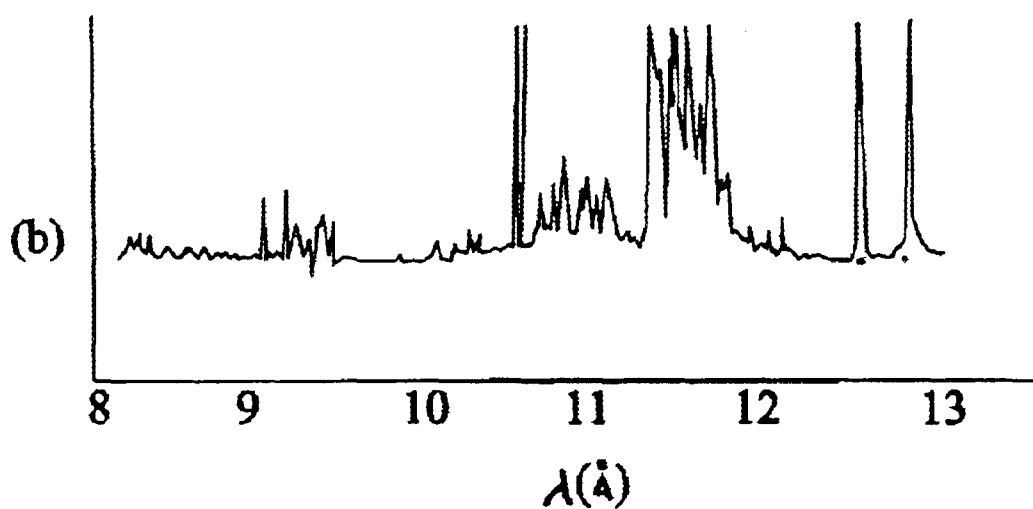
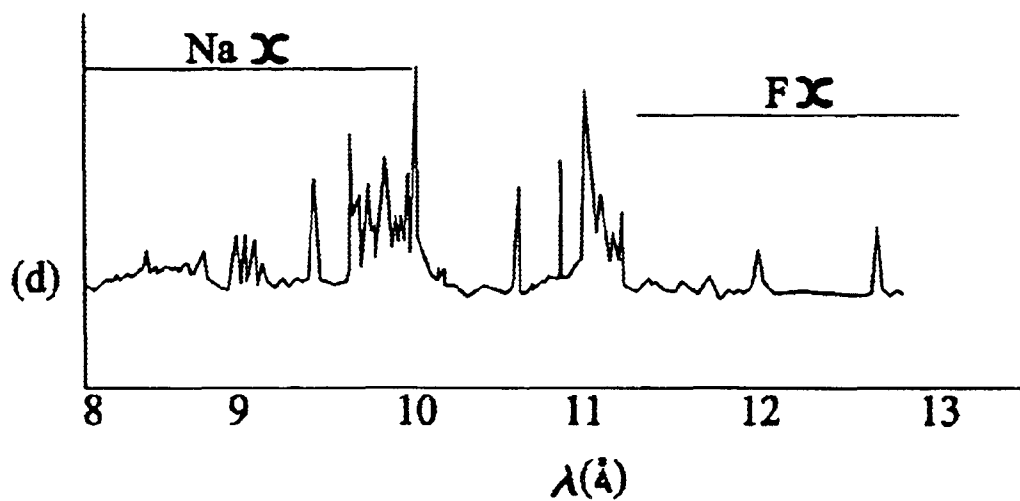


Fig.1b
(Prior Art)

Spectra of Zinc (Zn) target irradiated under similar
Scale : note 10Å = 1 nm



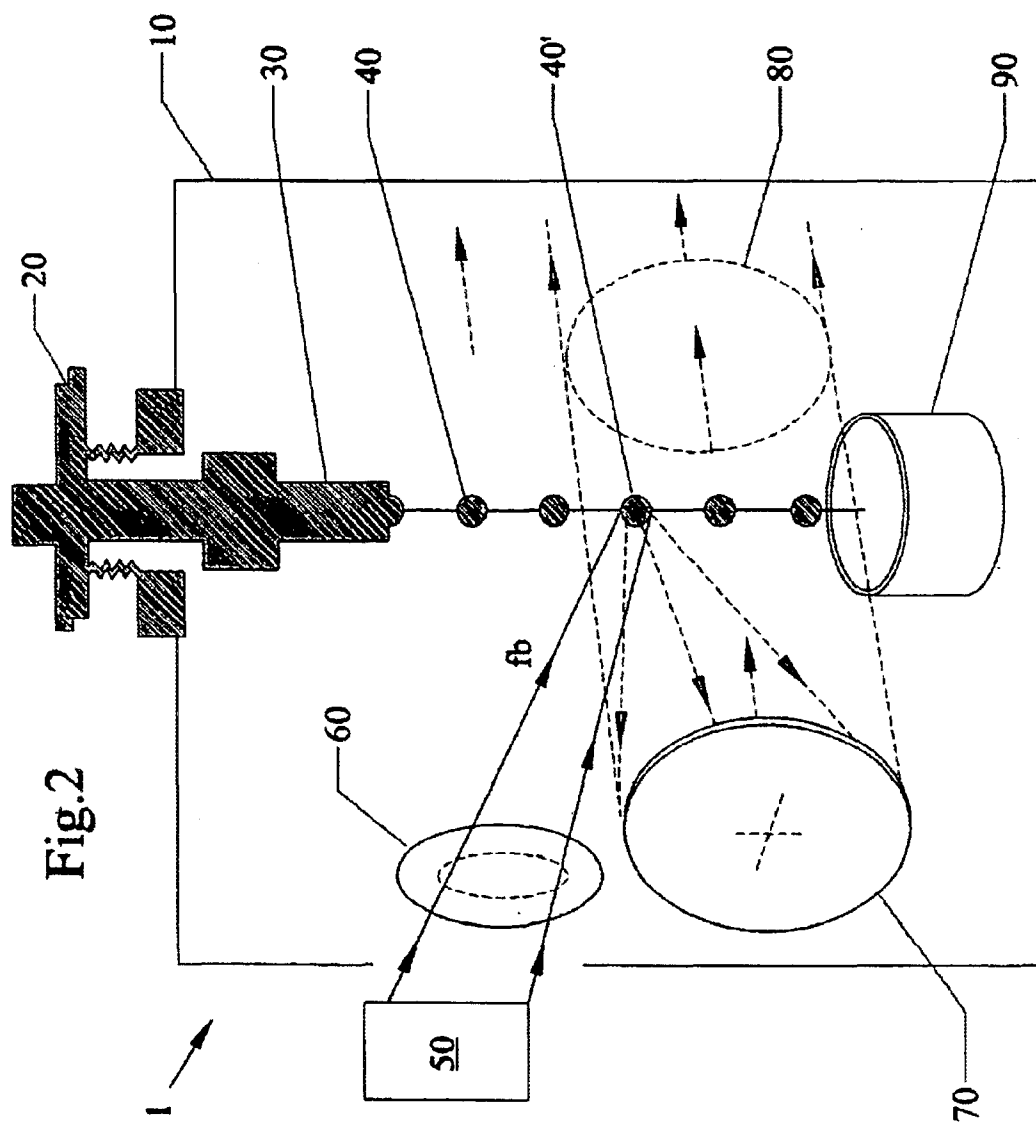


Fig.3a

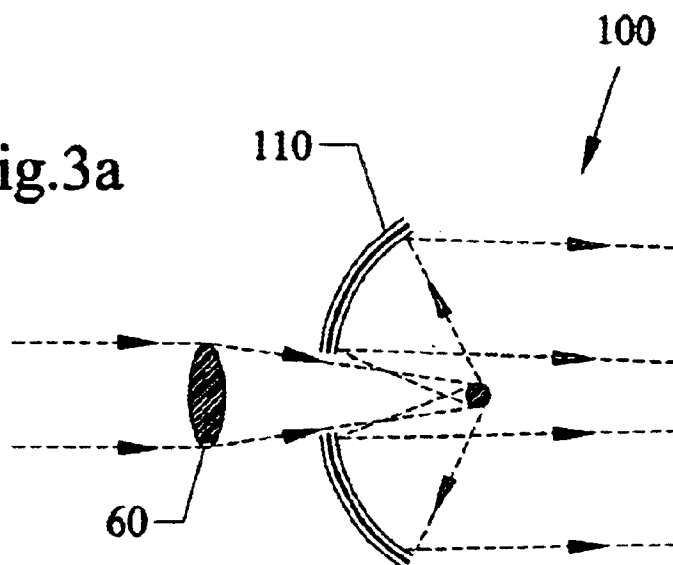


Fig.3b

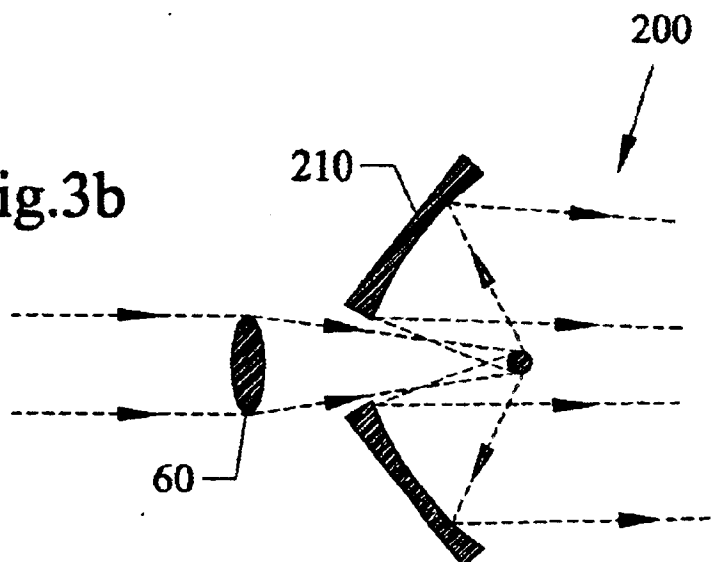
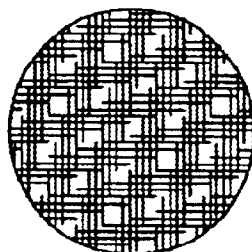


Fig.4



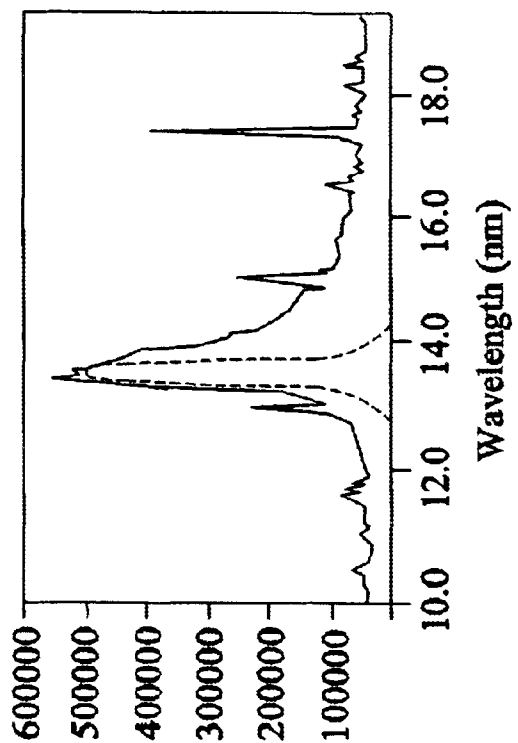


Fig. 5a

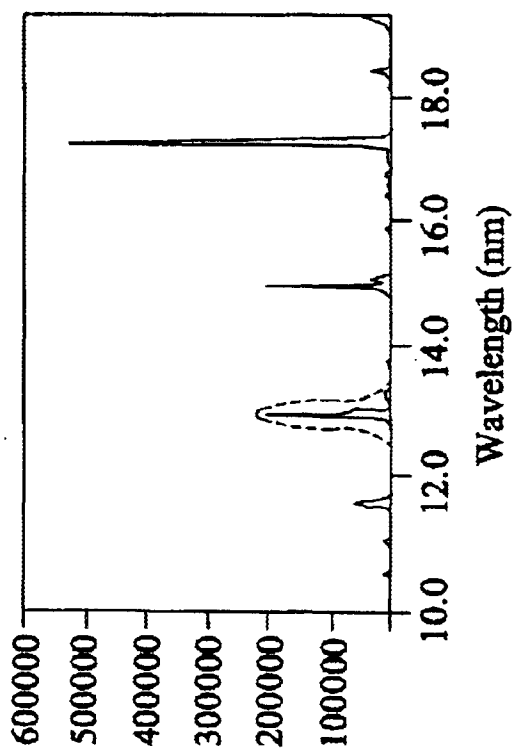


Fig. 5b

1

EUV, XUV, AND X-RAY WAVELENGTH SOURCES CREATED FROM LASER PLASMA PRODUCED FROM LIQUID METAL SOLUTIONS

This invention relates to laser point sources, and in particular to methods and apparatus for producing EUV, XUV and X-Ray type emissions from laser plasma produced from metal solutions being in liquid form at room temperature, and this invention claims the benefit of U.S. Provisional application No. 60/242,102 filed Oct. 20, 2000.

BACKGROUND AND PRIOR ART

The next generation lithographies (NGL) for advanced computer chip manufacturing have required the development of technologies such as extreme ultraviolet lithography (EUVL) as a potential solution. This lithographic approach generally relies on the use of multiplayer-coated reflective optics that has narrow pass bands in a spectral region where conventional transmissive optics is inoperable. Laser plasmas and electric discharge type plasmas are now considered prime candidate sources for the development of EUV. The requirements of this source, in output performance, stability and operational life are considered extremely stringent. At the present time, the wavelengths of choice are approximately 13 nm and 11.7 nm. This type of source must comprise a compact high repetition rate laser and a renewable target system that is capable of operating for prolonged periods of time. For example, a production line facility would require uninterrupted system operations of up to three months or more. That would require an uninterrupted operation for some 10 to the 9th shots, and would require the unit shot material costs to be in the vicinity of 10 to minus 6 so that a full size stepper can run at approximately 40 to approximately 80 wafer levels per hour. These operating parameters stretch the limitations of conventional laser plasma facilities.

Generally, laser plasmas are created by high power pulsed lasers, focused to micron dimensions onto various types of solids or quasi-solid targets, that all have inherent problems. For example, U.S. Pat. No. 5,151,928 to Hirose described the use of film type solid target tapes as a target source. However, these tape driven targets are difficult to construct, prone to breakage, costly and cumbersome to use and are known to produce low velocity debris that can damage optical components such as the mirrors that normally used in laser systems.

Other known solid target sources have included rotating wheels of solid materials such as Sn or tin or copper or gold, etc. However, similar and worse than to the tape targets, these solid materials have also been known to produce various ballistic particles sized debris that can emanate from the plasma in many directions that can seriously damage the laser system's optical components. Additionally these sources have a low conversion efficiency of laser light to in-band EUV light at only 1 to 3%.

Solid Zinc and Copper particles such as solid discs of compacted materials have also been reported for short wavelength optical emissions. See for example, T. P. Donaldson et al. Soft X-ray Spectroscopy of Laser-produced Plasmas, J. Physics, B:Atom. Molec. Phys., Vol. 9, No. 10, 1976, pages 1645-1655. FIGS. 1A and 1B show spectra emissions of solid Copper (Cu) and Zinc (Zn) targets respectively described in this reference. However, this reference requires the use of solid targets that have problems such as the generation of high velocity micro type projectiles that

2

causes damage to surrounding optics and components. For example, page 1649, lines 33-34, of this reference states that a "sheet of mylar . . . was placed between the lens and target in order to prevent damage from ejected target material . . ."

Thus, similar to the problems of the previously identified solids, solid Copper and solid Zinc targets also produce destructive debris when being used. Shields such as mylar, or other thin film protectors may be used to shield against debris for sources in the X-ray range, though at the expense of rigidity and source efficiency. However, such shields cannot be used at all at longer wavelengths in the XUV and EUV regions.

Frozen gases such as Krypton, Xenon and Argon have also been tried as target sources with very little success. Besides the exorbitant cost required for containment, these gases are considered quite expensive and would have a continuous high repetition rate that would cost significantly greater than \$ 10 to the minus 6. Additionally, the frozen gasses have been known to also produce destructive debris as well, and also have a low conversion efficiency factor.

An inventor of the subject invention previously developed water laser plasma point sources where frozen droplets of water became the target point sources. See U.S. Pat. Nos. 5,459,771 and 5,577,091 both to Richardson et al., which are both incorporated by reference. It was demonstrated in these patents that oxygen was a suitable emitter for line radiation at approximately 11.6 nm and approximately 13 nm. Here, the lateral size of the target was reduced down to the laser focus size, which minimized the amount of matter participating in the laser matter interaction process. The droplets are produced by a liquid droplet injector, which produces a stream of droplets that may freeze by evaporation in the vacuum chamber. Unused frozen droplets are collected by a cryogenic retrieval system, allowing reuse of the target material. However, this source displays a similar low conversion efficiency to other sources of less than approximately 1% so that the size and cost of the laser required for a full size 300 mm stepper running at approximately 40 to approximately 80 wafer levels per hour would be a considerable impediment.

Other proposed systems have included jet nozzles to form gas sprays having small sized particles contained therein, and jet liquids. See for Example, U.S. Pat. No. 6,002,744 to Hertz et al. and U.S. Pat. No. 5,991,360 to Matsui et al. However, these jets use many particles that are not well defined, and the use of jets creates other problems such as control and point source interaction efficiency. U.S. Pat. No. 5,577,092 to Kulak describe cluster target sources using rare expensive gases such as Xenon would be needed.

Attempts have been made to use a solid liquid target material as a series of discontinuous droplets. See U.S. Pat. No. 4,723,262 to Noda et al. However, this reference states that liquid target material is limited by example to single liquids such as "preferably mercury", abstract. Furthermore, Noda states that ". . . although mercury as been described as the preferred liquid metal target, any metal with a low melting point under 100 C. can be used as the liquid metal target provided an appropriate heating source is applied. Any one of the group of indium, gallium, cesium or potassium at an elevated temperature may be used . . .", column 6, lines 12-19. Thus, this patent again is limited to single metal materials and requires an "appropriate heating source (be) applied . . ." for materials other than mercury.

SUMMARY OF THE INVENTION

The primary objective of the subject invention is to provide an inexpensive and efficient target droplet system as

a laser plasma source for radiation emissions such as those in the EUV, XUV and x-ray spectrum.

The secondary objective of the subject invention is to provide a target source for radiation emissions such as those in the EUV, XUV and x-ray spectrum that are both debris free and that eliminates damage from target source debris.

The third objective of the subject invention is to provide a target source having an in-band conversion efficiency rate exceeding those of solid targets, frozen gasses and particle gasses, for radiation emissions such as those in the EUV, XUV and x-ray spectrum.

The fourth objective of the subject invention is to provide a target source for radiation emissions such as those in the EUV, XUV and x-ray spectrum, that uses metal liquids that do not require heating sources.

The fifth objective of the subject invention is to provide a target source for radiation emissions such as those in the EUV, XUV and x-ray spectrum that uses metals having a liquid form at room temperature.

The sixth objective of the subject invention is to provide a target source for radiation emissions such as those in the EUV, XUV and x-ray spectrum that uses metal solutions of liquids and not single metal liquids.

The seventh objective of the subject invention is to provide a target source for emitting plasma emissions at approximately 13 nm.

The eighth objective of the subject inventions is to provide a target source for emitting plasma emissions at approximately 11.6 nm.

The ninth objective of the subject invention is to provide a target source for x-ray emissions in the approximately 0.1 nm to approximately 100 nm spectral range.

A preferred embodiment of the invention uses compositions of metal solutions as efficient droplet point sources. The metal solutions include metallic solutions having a metal component where the metallic solution is in a liquid form at room temperature ranges of approximately 10 degrees C. to approximately 30 degrees C. The metallic solutions include molecular liquids or mixtures of elemental and molecular liquids. Each of the microscopic droplets of liquids of various metals with each of the droplets having diameters of approximately 10 micrometers to approximately 100 micrometers.

The molecular liquids or mixtures of elemental and molecular liquids can include a metallic chloride solution including ZnCl (zinc chloride), CuCl (copper chloride), SnCl (tin chloride), AlCl (aluminum chloride) and BiCl (bismuth chloride) and other chloride solutions. Additionally, the metal solutions can be a metallic bromide solutions such as CuBr, ZnBr, AlBr, or any other transition metal that can exist in a bromide solution at room temperature.

Other metal solutions can be made of the following materials in a liquid solvent. For example, Copper sulphate (CuSO₄), Zinc sulphate (ZnSO₄), Tin nitrate (SnSO₄), or any other transition metal that can exist as a sulphate can be used. Copper nitrate (CuNO₃), Zinc Nitrate (ZnNO₃), Tin nitrate (SnNO₃) or any other transition metal that can exist as a nitrate, can also be used.

Additionally, the metallic solutions can include organo-metallic solutions such as but not limited to CHBr₃ (Bromoform), CH₂I₂ (Diodomethane), and the like. Furthermore, miscellaneous metal solutions can be used such as but not limited to SeO₂ (38 gm/100 cc) (Selenium Dioxide), ZnBr₂ (447 gm/100 cc) (Zinc Dibromide), and the like.

Additionally, the metallic solutions can include mixtures of metallic nano-particles in liquids such as Al (aluminum) and liquids such as H₂O, oils, alcohols, and the like. Additionally, Bismuth and liquids such as H₂O, oils, alcohols, and the like.

The metallic solutions can be useful as target sources from emitting lasers that can produce plasma emissions at approximately 13 nm and approximately 11.6 nm.

Further objects and advantages of this invention will be apparent from the following detailed description of a presently preferred embodiment, which is illustrated schematically in the accompanying drawings.

BRIEF DESCRIPTION OF THE FIGURES

FIG. 1a shows a prior art spectra of using a solid Copper (Cu) target being irradiated.

FIG. 1b shows a prior art spectra of using Zinc (Zn) target being irradiated.

FIG. 2 shows a layout of an embodiment of the invention.

FIG. 3a shows a co-axial curved collecting mirror for use with the embodiment of FIG. 1.

FIG. 3b shows multiple EUV mirrors for use with embodiment of FIG. 1.

FIG. 4 is an enlarged droplet of a molecular liquid or mixture of elemental and molecular liquids that can be used in the preceding embodiment figures.

FIG. 5a is an EUV spectra of a water droplet target.

FIG. 5b is an EUV spectra of SnCl:H₂O droplet target (at approximately 23% solution).

DESCRIPTION OF THE PREFERRED EMBODIMENT

Before explaining the disclosed embodiment of the present invention in detail it is to be understood that the invention is not limited in its application to the details of the particular arrangement shown since the invention is capable of other embodiments. Also, the terminology used herein is for the purpose of description and not of limitation.

FIG. 2 shows a layout of an embodiment 1 of the invention. Vacuum chamber 10 can be made of aluminum, stainless steel, iron, or even solid-non-metallic material. The vacuum in chamber 10 can be any vacuum below which laser breakdown of the air does not occur (for example, less than approximately 1 Torr). The Precision Adjustment 20 of droplet can be a three axis position controller that can adjust the position of the droplet dispenser to high accuracy (micrometers) in three orthogonal dimensions. The droplet dispenser 30 can be a device similar to that described in U.S. Pat. Nos. 5,459,771 and 5,577,091 both to Richardson et al., and to the same assignee of the subject invention, both of which are incorporated by reference, that produces a continuous stream of droplets or single droplet on demand. Laser source 50 can be any pulsed laser whose focused intensity is high enough to vaporize the droplet and produce plasma from it. Lens 60 can be any focusing device that focuses the laser beam on to the droplet. Collector mirror 70 can be any EUV, XUV or x-ray optical component that collects the radiation from the point source plasma created from the plasma. For example it can be a normal incidence mirror (with or without multiplayer coating), a grazing incidence mirror, (with or without multiplayer coating), or some type of free-standing x-ray focusing device (zone plate, transmission grating, and the like). Label 90 refers to the EUV light which is collected. Cryogenic Trap 90 can be

5

a device that will collect unused target material, and possibly return this material for re-use in the target dispenser. Since many liquid targets used in the system will be frozen by passage through the vacuum system, this trap will be cooled to collect this material in the vacuum, until such time as it is removed. Maintaining this material in a frozen state will prevent the material from evaporating into the vacuum chamber and thereby increasing the background pressure. An increase in the background pressure can be detrimental to the laser-target interaction, and can serve to absorb some or all of the radiation produced by the plasma source. A simple configuration of a cryogenic trap, say for water-based targets, would be a cryogenically cooled "bucket" or container, into which the un-used droplets are sprayed. The droplets will stick to the sides of this container, and themselves, until removed from the vacuum chamber.

It is important that the laser beam be synchronized such that it interacts with a droplet when the latter passes through the focal zone of the laser beam. The trajectory of the droplets can be adjusted to coincide with the laser axis by the precision adjustment system. The timing of the laser pulse can be adjusted by electrical synchronization between the electrical triggering pulse of the laser and the electrical pulse driving the droplet dispenser. Droplet-on-demand operation can be effected by deploying a separate photodiode detector system that detects the droplet when it enters the focal zone of the laser, and then sends a triggering signal to fire the laser.

Referring to FIG. 2, after the droplet system 1 has been adjusted so that droplets are in the focal zone of the laser 50, the laser is fired. In high repetition mode, with the laser firing at rates of approximately 1 to approximately 100 kHz, the droplets or some of the droplets are plasmarized at 40'. EUV, XUV and/or x-rays 80 emitted from the small plasma can be collected by the collecting mirror 70 and transmitted out of the system. In the case where no collecting device is used, the light is transmitted directly out of the system.

FIG. 3a shows a co-axial curved collecting mirror 100 for use with FIG. 2. Mirror 110 can be a co-axial high NA EUV collecting mirror, such as a spherical, parabolic, ellipsoidal, hyperbolic reflecting mirror and the like. For example, like the reflector in a halogen lamp one mirror, axially symmetric or it could be asymmetric about the laser axis can be used. For EUV radiation it would be coated with a multi-layer coating (such as alternate layers of Molybdenum and Silicon) that act to constructively reflect light or particular wavelength (for example approximately 13 nm or approximately 1 nm or approximately 15 nm or approximately 17 nm, and the like). Radiation emanating from the laser-irradiated plasma source would be collected by this mirror and transmitted out of the system.

FIG. 3b shows multiple EUV mirrors for use with embodiment of FIG. 2. Mirrors 210 can be separate high NA EUV collecting mirrors such as curved, multilayer-coated mirrors, spherical mirrors, parabolic mirrors, ellipsoidal mirrors, and the like. Although, two mirrors are shown, but there could be less or more mirrors such as an array of mirrors depending on the application.

Mirror 210 of FIG. 3b, can be for example, like the reflector in a halogen lamp one mirror, axially symmetric or it could be asymmetric about the laser axis can be used. For EUV radiation it would be coated with a multi-layer coating (such as alternate layers of Molybdenum and Silicon) that act to constructively reflect light or particular wavelength (for example approximately 13 nm or approximately 11 nm or approximately 15 nm or approximately 17 nm, and the

6

like). Radiation emanating from the laser-irradiated plasma source would be collected by this mirror and transmitted out of the system.

FIG. 4 is an enlarged droplet of a metallic solution droplet. The various types of metal liquid droplets will be further defined in reference to Tables 1A–1F, which lists various metallic solutions that include a metal component that is in a liquid form at room temperature.

TABLE 1A

Metal chloride solutions	
ZnCl(zinc chloride)	
CuCl(copper chloride)	
SnCl(tin chloride)	
AlCl(aluminum chloride)	
Other transition metals that include chloride	

TABLE 1B

Metal bromide solutions	
CuBr (copper bromide)	
ZnBr (zinc bromide)	
SnBr (tin bromide)	
Other transition metals that can exist as a Bromide	

TABLE 1C

Metal Sulphate Solutions	
CuSO4 (copper sulphate)	
ZnSO4 (zinc sulphate)	
SnSO4 (tin sulphate)	
Other transition metals that can exist as a sulphate.	

TABLE 1D

Metal Nitrate Solutions	
CuNO3 (copper nitrate)	
ZnNO3 (zinc nitrate)	
SnNO3 (tin nitrate)	
Other transition metals that can exist as a nitrate	

TABLE 1E

Other metal solutions where the metal is in an organo-metallic solution.	
CHBr3(Bromoform)	
CH2I2(Diodomethane)	
Other metal solutions that can exist as an organo-metallic solution	

TABLE 1F

Miscellaneous Metal Solutions	
SeO2(38 gm/100 cc) (Selenium Dioxide)	
ZnBr2(447 gm/100 cc) (Zinc Dibromide)	

For all the solutions in Tables 1A–1F, the metal solutions can be in a solution form at a room temperature of approximately 10 degrees C. to approximately 30 degrees. Each of the droplet's diameters can be in the range of approximately 10 to approximately 100 microns, with the individual metal component diameter being in a diameter of that approaching approximately one atom diameter as in a chemical com-

pound. The targets would emit wavelengths in the EUV, XUV and X-ray regions.

FIG. 5a is an EUV spectrum of the emission from a pure water droplet target irradiated with a laser. It shows the characteristic lithium (Li) like oxygen emission lines with wavelengths at approximately 11.6 nm, approximately 13 nm, approximately 15 nm and approximately 17.4 nm. Other lines outside the range shown are also emitted.

FIG. 5b shows the spectrum of the emission from a water droplet seeded with approximately 25% solution of SnCl (tin chloride) irradiated under similar conditions. In addition to the Oxygen line emission, there is strong band of emission from excited ions of tin shown in the wavelength region of approximately 13 nm to approximately 15 nm. Strong emission in this region is of particular interest for application as a light source for EUV lithography. The spectrums for FIGS. 5a and 5b would teach the use of the other target solutions referenced in Tables 1A-1F.

As previously described, the novel invention is debris free because of the inherently mass limited nature of the droplet target. The droplet is of a mass such that the laser source completely ionizes (vaporizes) each droplet target, thereby eliminating the chance for the generation of particulate debris to be created. Additionally, the novel invention eliminates damage from target source debris, without having to use protective components such as but not limited to shields such as mylar or debris catchers, or the like.

Although the preferred embodiments describe individual tables of metallic type solutions, the invention can be practiced with combinations of these metallic type solutions as needed.

While the invention has been described, disclosed, illustrated and shown in various terms of certain embodiments or modifications which it has presumed in practice, the scope of the invention is not intended to be, nor should it be deemed to be, limited thereby and such other modifications or embodiments as may be suggested by the teachings herein are particularly reserved especially as they fall within the breadth and scope of the claims here appended.

I claim:

1. A method of producing optical emissions from a target source, comprising the steps of:

forming a metallic solution that includes molecular liquids or mixtures of elemental and molecular liquids at room temperature;

passing the metallic solution in microscopic droplets, each having a diameter of approximately 10 micrometers to approximately 100 micrometers into a target source; and

irradiating the target source with a high energy source to produce optical emissions that are debris free cannot cause debris damage to surrounding components.

2. The method of claim 1, wherein the high energy source includes; a laser source.

3. The method of claim 1, wherein the optical emissions include; X-rays.

4. The method of claim 1, wherein the optical emissions include: EUV (extreme ultraviolet) wavelength emissions.

5. The method of claim 1, wherein the optical emissions include: XUV wavelength emissions.

6. The method of claim 1, wherein the microscopic droplets each include: diameters of approximately 30 micrometers to approximately 90 micrometers.

7. The method of claim 6, wherein the microscopic droplets each include: diameters of approximately 40 micrometers to approximately 80 micrometers.

8. The method of claim 1, wherein the metallic solution includes: a metallic chloride solution.

9. The method of claim 8, wherein the metallic chloride solution includes: ZnCl(zinc chloride).

10. The method of claim 8, wherein the metallic chloride solution includes: CuCl(copper chloride).

11. The method of claim 8, wherein the metallic chloride solution includes: SnCl(tin chloride).

12. The method of claim 8, wherein the metallic chloride solution includes: AlCl (aluminum chloride).

13. The method of claim 1, wherein the metallic solution includes:

a metallic bromide solution.

14. The method of claim 13, wherein the metallic bromide solution includes:

CuBr(copper bromide).

15. The method of claim 13, wherein the metallic bromide solution includes:

ZnBr(zinc bromide).

16. The method of claim 13, wherein the metallic bromide solution includes:

SnBr(tin bromide).

17. The method of claim 1, wherein the metallic solution includes:

a metallic sulphate solution.

18. The method of claim 17, wherein the metallic sulphate solution includes:

CuSO4(copper sulphate).

19. The method of claim 17, wherein the metallic sulphate solution includes:

ZnSO4(zinc sulphate).

20. The method of claim 17, wherein the metallic sulphate solution includes:

SnSO4(tin sulphate).

21. The method of claim 1, wherein the metallic solution includes:

a metallic nitrate solution.

22. The method of claim 21, wherein the metallic nitrate solution includes:

CuNO3(copper nitrate).

23. The method of claim 21, wherein the metallic nitrate solution includes:

ZnNO3(zinc nitrate).

24. The method of claim 21, wherein the metallic nitrate solution includes:

SnNO3(tin nitrate).

25. The method of claim 1, wherein the room temperature includes:

approximately 10 degrees C. to approximately 30 degrees C.

26. The method of claim 1, wherein the optical emissions include:

approximately 11.7 nm.

27. The method of claim 1, wherein the optical emissions include:

approximately 13 nm.

28. The method of claim 1, wherein the metallic solution includes:

an organo-metallic solution.

29. The method of claim 28, wherein the organo-metallic solution includes:

CHBr3(Bromoform).

9

30. The method of claim 28, wherein the organo-metallic solution includes:

CH₂I₂(Diodomethane).

31. The method of claim 1, wherein the metallic solution includes:

SeO₂(Selenium Dioxide).

32. The method of claim 1, wherein the metallic solution includes:

ZnBr₂ (Zinc Dibromide).

33. An method of generating optical emissions from metallic point sources, comprising the steps of:

forming microscopic liquid metal droplets at room temperature without heating the droplets;

passing the droplets, each having a diameter in the range of approximately 10 to approximately 100 microns, into individual target sources;

irradiating the individual target sources with a laser beam having substantially identical diameter to each of the individual droplets; and

producing optical emissions from the irradiated target sources without debris damage to surrounding components.

34. The method of claim 33, wherein each of the microscopic liquid metal droplets include: metallic chloride solutions.

35. The method of claim 33, wherein each of the microscopic liquid metal droplets include: metallic bromide solutions.

36. The method of claim 33, wherein each of the microscopic liquid metal droplets include: metallic sulphate solutions.

37. The method of claim 33, wherein each of the microscopic liquid metal droplets include: metallic nitrate solutions.

38. The method of claim 33, wherein each of the microscopic liquid metal droplets include: an organo-metallic solution.

39. The method of claim 33, wherein the room temperature includes:

approximately 10 degrees to approximately 30 degrees C.

40. The method of claim 33, wherein the optical emissions include:

approximately 11.7 nm.

41. The method of claim 33, wherein the optical emissions include:

approximately 13 nm.

42. The method of claim 34, wherein the metallic chloride solution

includes: ZnCl₂(zinc chloride).

43. The method of claim 34, wherein the metallic chloride solution

includes: CuCl₂(copper chloride).

44. The method of claim 34, wherein the metallic chloride solution

includes: SnCl₂(tin chloride).

45. The method of claim 33, wherein each of the microscopic liquid metal droplets include: approximately 25% metallic solutions.

46. A method of producing optical emissions from liquid droplet target sources, comprising the steps of:

forming liquid metal droplets at room temperature;

passing the liquid metal droplets into individual target sources; and

irradiating the target sources with a high energy source to produce optical emissions that are debris free and cannot cause debris damage to surrounding components.

10

47. The method of claim 46, wherein each of the target source droplets include approximately 25% metallic solutions.

48. The method of claim 47, wherein each of the droplets are microscopic with a diameter of approximately 10 micrometers to approximately 100 micrometers.

49. The method of claim 48, wherein the diameters of the droplets are approximately 30 micrometers to approximately 90 micrometers.

50. The method of claim 48, wherein the diameters of the droplets are approximately 4 micrometers to approximately 80 micrometers.

51. The method of claim 46, wherein each of the liquid metal

droplets include: metallic chloride solutions.

52. The method of claim 46, wherein each of the liquid metal

droplets include: metallic bromide solutions.

53. The method of claim 46, wherein each of the liquid metal

droplets include: metallic sulphate solutions.

54. The method of claim 46, wherein each of the liquid metal

droplets include: metallic nitrate solutions.

55. The method of claim 46, wherein each of the liquid metal

droplets include: an organo-metallic solution.

56. The method of claim 46, wherein the room temperature includes:

approximately 10 degrees to approximately 30 degrees C.

57. The method of claim 51, wherein the metallic chloride solutions

includes: ZnCl₂(zinc chloride).

58. The method of claim 51, wherein the metallic chloride solutions

includes: CuCl₂(copper chloride).

59. The method of claim 51, wherein the metallic chloride solutions

includes: SnCl₂(tin chloride).

60. An apparatus for generating optical emissions from liquid point sources, comprising:

means for forming liquid metal droplets at room temperature;

means for feeding the liquid metal droplets at room temperature into a target path to form individual target sources;

means for irradiating the individual target sources with an optical beam; and

means for generating optical emissions from the irradiated target sources that are debris free and cannot cause debris damage to surrounding components.

61. The apparatus of claim 60, wherein the irradiating means includes: a laser.

62. The apparatus of claim 60, wherein each of the liquid metal droplets are microscopic sized droplets have a diameter of approximately 10 micrometers to approximately 100 micrometers.

63. The apparatus of claim 62, wherein the diameters of each of the liquid metal droplets are approximately 30 micrometers to approximately 90 micrometers.

64. The apparatus of claim 62, wherein the diameters of each of the liquid metal droplets are approximately 40 micrometers to approximately 80 micrometers.

11

65. The apparatus of claim 60, wherein the target sources include:

approximately 25% metallic solutions.

66. The apparatus of claim 60, wherein each of the liquid metal droplets include: metallic chloride solutions.

67. The apparatus of claim 60, wherein each of the liquid metal droplets include: metallic bromide solutions.

68. The apparatus of claim 60, wherein each of the liquid metal droplets include: metallic sulphate solutions.

69. The apparatus of claim 60, wherein each of the liquid metal droplets include: metallic nitrate solutions.

70. The apparatus of claim 60, wherein each of the liquid metal droplets include: organo-metallic olutions.

12

71. The apparatus of claim 60, wherein the room temperature includes: approximately 10 degrees to approximately 30 degrees C.

72. The apparatus of claim 66, wherein the metallic chloride solutions includes: ZnCl(zinc chloride).

73. The apparatus of claim 66, wherein the metallic chloride solutions includes: CuCl(copper chloride).

74. The method of claim 66, wherein the metallic chloride solutions includes: SnCl(tin chloride).

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